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1 **Occurrence and assemblage composition of intertidal non-native species may be** 2 **influenced by shipping patterns and artificial structures**

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17 18 **Abstract**

19 Habitat modification coupled with the spread of non-native species (NNS) are among the top
20 threats to marine biodiversity globally. Species are known to be transported to new locations
21 via international shipping and secondarily spread via regional vessels and artificial structures.
22 Rapid Assessment Surveys (RAS) combining quantitative and semi-quantitative methods
23 compared NNS richness and assemblage composition on intertidal natural rocky shores and
24 artificial structures in harbours in different regions along the south coast of England.
25 Quantitative data showed that artificial habitats supported higher richness than natural
26 habitats, while semi-quantitative data found no difference in richness among habitat types.

This result was attributed to additional species found in rock pools during searches of complex microhabitats in natural habitats. Assemblages on artificial structures differed among regions, with regions and harbours with greater numbers of vessels supporting greater richness. Results highlight the importance of shipping and artificial structures for NNS introduction and spread.

Keywords

Invasive species, non-indigenous species, biodiversity, biological invasion, ocean sprawl, Rapid Assessment Surveys

1. Introduction

Habitat modification and the introduction and spread of non-native species (NNS) are impacting natural ecosystems and threatening global biodiversity (Manchester and Bullock, 2000; Bax et al., 2003; Simberloff, 2005). In marine coastal systems, “ocean sprawl” (*sensu* Duarte et al., 2012) – the proliferation of artificial structures (e.g., seawalls, groynes, piers, floating pontoons, offshore platforms) – is replacing natural habitats with a variety of hard engineered structures built to support human activities (e.g., aquaculture, transportation, industry, shipping, energy extraction), as well as stabilise and protect shorelines from rising and stormier seas (Griggs, 2005; Duarte et al., 2012; Firth et al., 2016a; Bishop et al., 2017). Artificial structures provide new ‘competitor-free’ habitat for NNS settlement and establishment (Airoldi and Bulleri, 2011; Firth et al., in review), as well as increase ecological connectivity between local and global shipping hubs (Floerl et al., 2009; Airoldi et al., 2015). These novel habitats enable the spread of cryptogenic (i.e., it is unclear whether the species is native or introduced; Kinzie, 1984; Carlton, 1996a), opportunistic (i.e., a species adapted to exploit new or disturbed habitats; Whitlatch and Zajac, 1985) and non-native species (Ruiz et al., 1997;

Dafforn et al., 2009; Firth et al., 2016a). Artificial structures also facilitate the homogenisation of biological communities, supporting novel species assemblages not encountered in natural habitats, and affect the structure and functioning (physical and ecological) of the surrounding environment (McKinney and Lockwood, 1999; McKinney, 2006; Dugan et al., 2011).

The rise in global trade has meant that harbours are characterised by a range of artificial structures, with an extraordinary amount of shipping traffic arriving from ports across the globe every day (Seebens et al., 2013; Marine Traffic, 2019; World Port Source, 2019). These mobile vectors (ranging from small, local leisure craft to large inter-continental commercial tankers) are able to spread NNS among the proliferating static structures (i.e., seawalls, breakwaters, groynes, floating pontoons) in destination ports such that the latter act as species reservoirs (Ruiz et al., 1997; Neves et al., 2007; Clarke Murray et al., 2011; Mineur et al., 2012). At a global scale, the primary vectors of initial introduction are typically transoceanic ships, barges and floating platforms (i.e., mobile vectors) that dock in large international harbours (Carlton and Geller, 1993; Ruiz et al., 1997; Molnar et al., 2008). These vessels transport species in two main ways: (1) as larvae in ship ballast water (Ruiz et al., 1997; Gollasch, 2008) and (2) as adults fouling ship hulls (Gollasch, 2002; Drake and Lodge, 2007). Thus, initial NNS colonisation and settlement tend to be highest within major shipping ports compared to surrounding areas (Eno et al., 1997; Molnar et al., 2008; Keller et al., 2011). Secondary, local spread of NNS is then probably through a combination of small mobile vectors (fishing and leisure craft) to nearby artificial structures (Clarke Murray et al., 2011; Mineur et al., 2012; Airoidi et al., 2015). Planktonic larvae can also be carried away from the port of introduction by wave-driven currents (McQuaid and Phillips, 2000), settling on artificial structures along the coast (McQuaid and Phillips, 2000; Wasson et al., 2001). In this way, artificial structures can act as stepping stones, allowing non-natives to persist or spread by provision of 'virgin' hard substrate amongst otherwise uninhabitable habitats (e.g., 'soft bottom' sediment habitat; Apte et al., 2000; Sammarco, 2015; Airoidi et al., 2015).

78 The biological communities of artificial structures are typically less diverse, and support
79 greater numbers of NNS, than comparable, nearby natural rocky shore habitats (Glasby et al.,
80 2007; Vaselli et al., 2008; Dafforn et al., 2015a). This disparity is attributed to the physical
81 design of artificial structures; they typically have steep profiles and reduced surface area and
82 limited topographic complexity compared to their natural analogues (Moschella et al., 2005;
83 Chapman and Underwood, 2011). The disturbance associated with human activities in
84 harbours can also physically dislodge organisms and create space for new colonisers to
85 exploit, thereby influencing successional dynamics of the community (e.g., removal of
86 predators, loss of canopy algae; Stachowicz et al., 1999; Byers, 2002). Additionally, ports are
87 usually located within sheltered bays or estuaries, which by nature, experience greater
88 fluctuations in temperature and salinity (Whitehead et al., 2009), input of nutrients (Statham,
89 2012) and other pollutants (Stark, 1998; Johnston et al., 2017; Hitchcock and Mitrovic, 2019)
90 compared to open coasts. More importantly, many non-natives are generalist species that
91 often have longer planktonic larval durations or extended reproductive seasons (Dineen et al.,
92 2001; Muxagata et al., 2004), which means they are able to take advantage of bare space as
93 it becomes available through creation of new substrate or after disturbance events. For
94 example, in the UK, the non-native barnacle, *Austrominius modestus* (Darwin, 1854), is
95 reproductive almost year-round (Muxagata et al., 2004), while native barnacle species
96 reproduce mainly in the spring (e.g., *Semibalanus balanoides* (Linnaeus, 1767)) or summer
97 (e.g., *Chthamalus montagui* Southward, 1976 and *C. stellatus* (Poli, 1791); Burrows et al.,
98 1992). These physical and biological factors probably interact, leaving severely disturbed
99 areas vulnerable to more resilient and opportunistic invaders (Stachowicz et al., 1999; Airolidi
100 and Bulleri, 2011; Johnston et al., 2017).

101 It is important to understand the practically synergistic interaction between ocean sprawl
102 and global shipping so that potential introduction points can be predicted and appropriate

invasive species forecasting techniques can be developed. Yet to our knowledge, we know of no studies that explored the influence of shipping patterns on occurrence of NNS in intertidal habitats along the south coast of England (an area historically known as a point of introduction to the British Isles; for a review of potential sources of NNS introduction into the British Isles see Eno et al. (1997)). Even less attention has focussed on understanding differences in NNS occurrence and assemblage compositions between natural and artificial intertidal habitats (but see Glasby et al., 2007; Dafforn et al., 2012; Dafforn et al., 2015a for natural and artificial comparison in subtidal habitats). Information from natural habitats may be useful in determining the potential for NNS to spread out from points of initial introduction (Valentine et al., 2007; Carman and Grunden, 2010; Epstein and Smale, 2018). Understanding the mechanisms underpinning the differences in NNS occurrence between natural and artificial habitats is also critical to develop a robust foundation of evidence upon which to base ecological engineering (i.e., the combination of ecological and engineering design to create sustainable ecosystems that benefit humans and nature; Mitsch and Jørgensen, 2003; Odum and Odum, 2003). To address the current knowledge gaps concerning occurrence of NNS on natural rocky shores and artificial structures in intertidal habitats, we conducted surveys of NNS in intertidal natural and artificial sites in 11 harbours along the south coast of England to test the following hypotheses:

1. NNS richness would be greater, and NNS assemblage composition would differ, in artificial compared to natural habitats.
2. NNS richness and NNS assemblage composition in artificial habitats would differ among harbours and regions, but would be highest in ports with higher number of arriving vessels.

Our study also provided the opportunity to compare data obtained from quantitative stratified-random quadrat-based surveys with semi-quantitative methods based on timed searches, as both approaches were used here.

2. Materials and methods

2.1. Study region

The English Channel is one of the busiest waterways in the world (Marine Traffic, 2019; World Port Source, 2019). The coast along the English side of the Channel is characterised by a number of harbours that support international and regional shipping and cruise traffic, military traffic, ferries to continental Europe and recreational and tourist activities (Table 1). Consequently, the south coast of England has traditionally been susceptible to invasions and is known as the point of introduction into the British Isles for many NNS from Europe and around the world (Eno et al., 1997; Bishop et al., 2015a; Bishop et al., 2015b). In this study, harbours were grouped into geographic regions following Bishop et al. (2015b) (West, Central, East; Table 1, Figure 1). One survey was done per site (hereafter referred to as 'site'), and sites were located either within natural (rocky shores) or artificial (artificial structures) habitats. Numbers of sites surveyed per harbour varied and reflected the size of the harbour. All harbours had artificial substrata, but only harbours in the West had natural rocky shore (contained within the larger natural harbour) for comparison with artificial structures (for classification of harbours within regions see Table 1). The natural versus artificial habitat assessment involved only seawalls made from naturally-sourced rock for comparison to natural rocky shores. The comparison of NNS across all harbours along the south coast of England assessed multiple types of artificial structures, which included piers, marina wave-breaker walls, seawalls, discharge pipes, groynes, boat docks, bridge support structures, wharfs and breakwaters (Table S1). As artificial structures were opportunistically sampled, not all structure types were represented in each harbour. To compare richness and assemblage composition of artificial sites (artificial structures) among harbours and regions across the entire south coast of England, all artificial structures were used for analysis. The areas

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154 surveyed were at the mouths of any estuarine complexes and generally fully saline at high
155 tides and thus comparable to non-estuarine ports (i.e., Torbay, Folkestone, Dover; Table S1).

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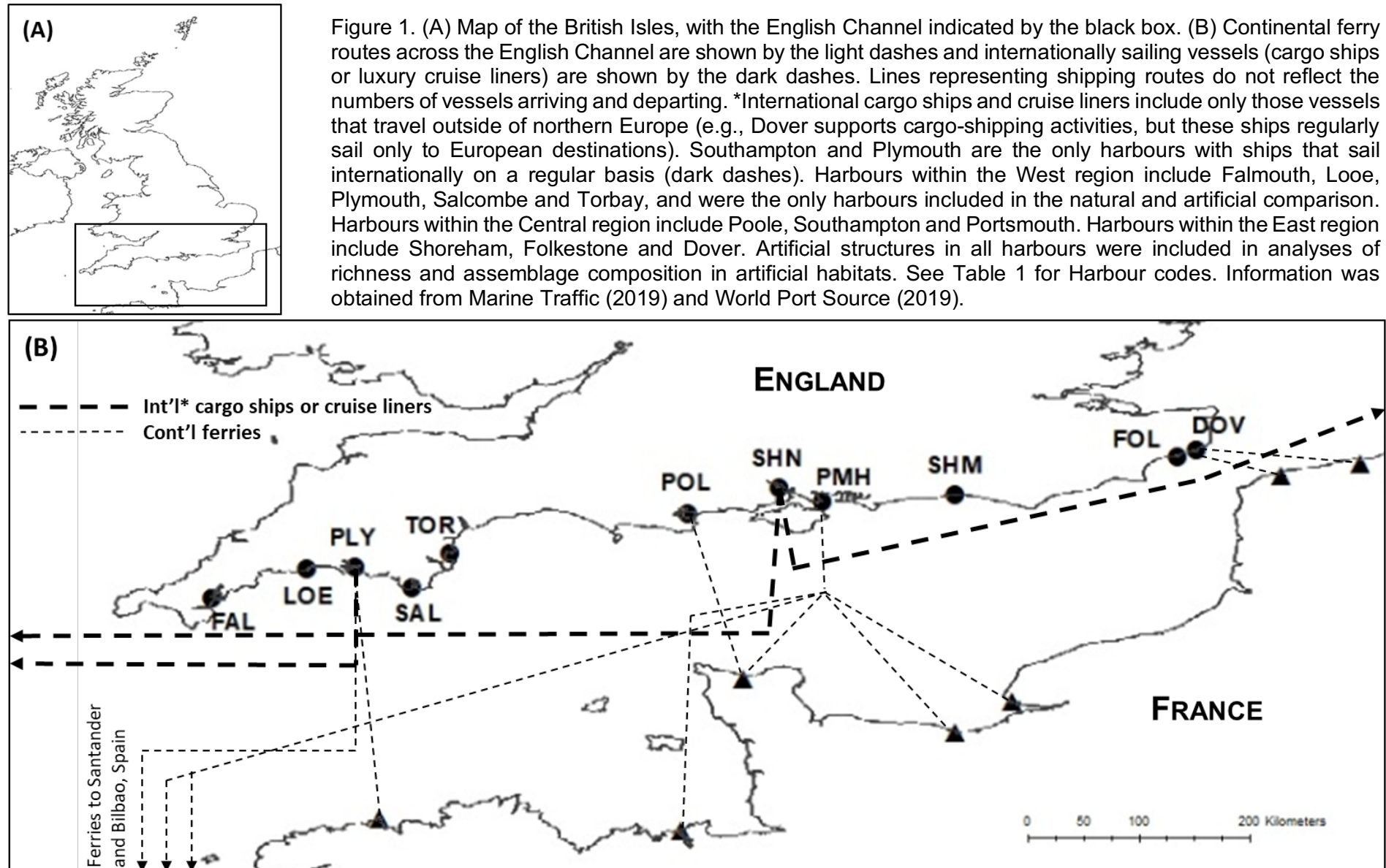
160 Table 1. Details for NNS RAS conducted along the south coast of England, including survey details, type of vessels by harbour and harbour
161 characteristics. Natural and artificial habitat comparisons were only done in the West region (FAL, LOE, PLY, SAL, TOR). Vessel information was
162 obtained from Marine Traffic (2019) and World Port Source (2019). *International shipping describes types of destinations. **Continental
163 passenger ferries (number of destinations) travel from south England to northern Europe. †International cruise lines describe types of destinations.
164 ‡Dominant natural habitat is rocky shore ('RS') or soft bottom ('SB') habitat. ††Main features include ^asize of harbour ('v. sm' = very small, 'sm' =
165 small, 'med' = medium and 'lg' = large); ^btype of harbour (natural coastal inlet, coastal breakwater); ^cfreshwater input; ^ddepth of main channel
166 ('shallow' = < 5 m, 'average' = 5-9 m and 'deep' = > 9 m) and ^eaverage tidal range as recorded in July 2018 from Tide Plotter (v. 5.8, Belfield
167 Software Ltd). Information on size and type of harbour, as well as depth of main channel were obtained from World Port Source (2019). Details
168 regarding type of artificial structures surveyed in each harbour is provided in Table S1.

Survey details			Type of vessels					Harbour characteristics	
Har code	Harbour/city	Region	Int'l shipping*	Cont'l passenger ferry **	Int'l cruise lines†	Military	Fishing vessels & leisure craft	Dom nat hab‡	Main features
FAL	Falmouth	West	0	0	0	✓	✓	RS	med ^a ; natural coastal inlet ^b ; moderate input ^c ; average ^d ; 3.6 m ^e
LOE	Looe	West	0	0	0	0	✓	RS	v. sm ^a ; natural coastal inlet ^b ; major input ^c ; shallow ^d ; 3.5 m ^e
PLY	Plymouth	West	Global	2	0	✓	✓	RS	med ^a ; natural coastal inlet ^b ; moderate input ^c ; deep ^d ; 3.6 m ^e
SAL	Salcombe	West	0	0	0	0	✓	RS	v. sm ^a ; natural coastal inlet ^b ; minimal input ^c ; shallow ^d ; 3.3 m ^e
TOR	Torbay	West	0	0	0	0	✓	RS	v. sm ^a ; coastal breakwater ^b ; no input ^c ; average ^d ; 3.0 m ^e
POL	Poole	Central	0	1	0	0	✓	SB	sm ^a ; natural coastal inlet ^b ; limited input ^c ; shallow ^d ; 1.0 m ^e
SHN	Southampton	Central	Global	0	Global	0	✓	SB	lg ^a ; natural coastal inlet ^b ; moderate input ^c ; average ^d ; 2.9 m ^e
PMH	Portsmouth	Central	0	6	0	✓	✓	SB	med ^a ; natural coastal inlet ^b ; limited input ^c ; average ^d ; 3.1 m ^e
SHM	Shoreham	East	0	0	0	0	✓	SB	v. sm ^a ; natural coastal inlet ^b ; major input ^c ; shallow ^d ; 4.5 m ^e
FOL	Folkestone	East	0	0	0	0	✓	SB	v. sm ^a ; coastal breakwater ^b ; no input ^c ; average ^d ; 5.2 m ^e
DOV	Dover	East	European	2	European	0	✓	SB	sm ^a ; coastal breakwater ^b ; no input ^c ; average ^d ; 4.7 m ^e

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2.2. Sampling methods

Rapid Assessment Surveys (RAS) provide a pragmatic method of covering a large number of locations in a reasonable amount of time (Pederson et al., 2005; Arenas et al., 2006; Bishop et al., 2015a,b); they typically involve a qualitative approach, using timed searches of targeted areas and habitats (e.g., undersides of floating pontoons). In biodiversity surveys, however, the most common means of gathering data is predominantly through quantitative quadrat sampling, often avoiding topographically complex surfaces (Bulleri et al., 2005; Dafforn et al., 2012; Firth et al., 2016b). In this study, a combined approach was employed to capture NNS richness and abundance (i.e., diversity), which consisted of utilising quantitative stratified-random quadrat sampling and semi-quantitative timed searches. All artificial structures were accessed on foot at low tide. Native biota were not quantified during any of the surveys. Quantitative stratified-random quadrat sampling involved haphazardly placing 20 quadrats (25 x 25 cm) in the lower intertidal within a 10 x 10 m area and recording counts of mobile organisms and percentage cover of sessile organisms. For the purpose of this study, the lower intertidal was the area of the shore that was inundated during neap low tides but exposed at spring low tides (i.e., surveys occurred only on spring low tides when tide was ≤ 1 m above CD). Occasionally, this area was condensed because the steeper slope of artificial structures resulted in reduced area available to survey. In these cases, a longer horizontal section was sampled to compensate for lost vertical area. All NNS visible to the naked eye within the quadrats were identified and quantified. To positively identify and quantify the non-native barnacle, *Austrominius modestus*, 5 x 5 cm photo-quadrat images ($n = 20$) were taken in the densest barnacle zone and photographs were later analysed using ImageJ (Schneider et al., 2012). Slope and substrate were standardised by surveying vertical or sloping substrate ($\geq 45^\circ$ angle) and avoiding topographically complex surfaces (i.e., gaps, grooves, pits, crevices, rock pools). To locate rare species, one person conducted a 30-minute timed search across

the study area, including complex surfaces and microhabitats (e.g., crevices, rock pools, undersides of boulders). Additional minutes were added to the search where logistical constraints delayed efficient sampling (e.g., some artificial structures had characteristics that required careful manoeuvring around the structure compared to other easily accessible structures or natural rocky shore). A semi-quantitative search-based assessment of overall abundance of each NNS was made on a scale of 0-3 (0 = absent, 1 = rare-occasional, 2 = frequent-common, 3 = abundant-superabundant; Bishop et al., 2015b). Thus, the quantitative quadrat method produced 20 quadrat replicates per site, while the semi-quantitative search-based method produced one abundance score for each NNS per site. Species that could not be identified in the field (e.g., bryozoans such as *Tricellaria inopinata* (d'Hondt & Occhipinti Ambrogi, 1985) and *Bugulina* spp.) were preserved in 70% ethanol and transported back to the laboratory for microscopic examination.

2.2.1. Comparison of NNS in natural and artificial habitats

To investigate differences of occurrence of NNS between natural and artificial habitats, NNS richness and assemblage composition were recorded in ten natural sites (rocky shores) and eleven artificial sites (seawalls) in the West region (Table 1). Natural sites were chosen based on location to closest harbour and were as sheltered as possible to reduce the influence of wave exposure gradients on assemblage composition. There was no restriction placed on size of seawall.

2.2.2. Comparison of NNS in artificial habitats along the south coast of England

To assess differences of occurrence of NNS in artificial habitats among harbours and regions, eleven harbours spanning three regions across the entire south coast of England were surveyed. Harbours within regions were chosen based on major harbours surveyed in Bishop et al. (2015b) and those which stretched across each region. As many artificial sites as possible with public access were surveyed in each harbour (Table 1).

2.3. Statistical analyses

2.3.1. Comparison of NNS in natural and artificial habitats

To compare NNS richness and assemblage composition between natural and artificial habitats, comparisons were made between ten natural sites and eleven nearby artificial sites using both quantitative quadrat and semi-quantitative search-based data separately (i.e., data from both methods were used but were analysed separately). For both sampling methods, where abundance information was used, data were fourth-root transformed to down-weight the influence of very abundant species (Anderson et al., 2008). Bray-Curtis dissimilarity matrices were then computed, and permutational multivariate analysis of variance tests (PERMANOVA; Anderson, 2001) were used to test for differences in species richness and assemblage composition. Two-way nested designs with Habitat as a fixed factor (2 levels: natural, artificial) and Site as a random factor (nested in Habitat) were employed. PERMANOVA tests were based on 9999 permutations of residuals under a reduced model. Tests for differences were conducted in PRIMER v6 with the PERMANOVA+ add-on using the PERMANOVA routine (PRIMER-E Ltd, Plymouth, UK; Anderson et al., 2008). Ordination of samples were visualised using two-dimensional non-metric multidimensional scaling (nMDS) plots. Contributions to dissimilarities among regions from each species was determined using the similarity percentages routine (SIMPER).

2.3.2. Comparison of NNS in artificial habitats along the south coast of England

Differences in NNS richness and assemblage composition in artificial habitats among harbours and regions were assessed using data from both the quantitative quadrat and semi-quantitative search-based surveys. Where abundance information was used, data were fourth-root transformed. Bray-Curtis dissimilarity matrices were then computed, and permutational multivariate analysis of variance tests (PERMANOVA) were used to test for differences in species richness and assemblage composition. For quantitative quadrat data, a

three-way nested design was used for each test (species richness and assemblage composition) with Site as a random factor (number of levels depended on number of sites surveyed in each harbour) nested in Harbour as a fixed factor (number of levels varied with region) nested in Region as a fixed factor (3 levels: West, Central, East). As quantitative quadrat sampling was not undertaken in Salcombe, only ten harbours were included in quantitative quadrat analyses. Because there was not replication at the 'Site' level when semi-quantitative search-based data were analysed (i.e., there was one abundance value per site), two-way nested designs were used with Harbour as a fixed factor (number of levels varied with region) nested in Region as a fixed factor (3 levels) for each test (species richness and assemblage composition). Information about vessel type and the average number of vessels per harbour over a 60-day period as a proxy for boat traffic in general was obtained from the Marine Traffic (Marine Traffic, 2019) and World Port Source (World Port Source, 2019) websites. General observations comparing numbers of NNS with numbers and types of vessels were made with no formal analyses done.

3. Results

3.1. Overall results

A total of 26 NNS were recorded from natural and artificial habitats across the entire south coast of England (Table S1). Fifteen NNS were recorded from the natural and artificial habitat comparison in the West region (Falmouth to Torbay); 12 of these NNS were found in artificial, while 9 NNS were recorded in natural habitats (Table S1). Six species were exclusive to artificial habitats, while 3 species were exclusive to natural habitats, with 6 species common to both. Two NNS were discovered in new localities: colonies of the carpet sea squirt, *Didemnum vexillum* Kott, 2002, were found on the seaward side of a wooden wave-breaker wall and a metal pipe positioned perpendicular to the shore in Poole Harbour. Colonies were

also found in Portsmouth on concrete fishing piers perpendicular to the shore. The red alga, *Botryocladia wrightii* (Harvey) W.E.Schmidt, D.L.Ballantine & Fredericq, 2017 (recently changed from *Chrysomenia wrightii*), was found in a small water-retaining pool along a stepped seawall in Portsmouth. The only previously confirmed records of *B. wrightii* in this study region were from marinas in Falmouth (Wood et al., 2015).

3.2. Comparison of NNS in natural and artificial habitats

Of the 15 NNS recorded across natural and artificial habitats between Falmouth and Torbay (Table S1), 9 taxa were recorded in natural (60% of total), while 12 were observed in artificial (80% of total) habitats. Species unique to natural habitats included the brown alga, *Undaria pinnatifida* (Harvey) Suringar, 1873 and the red algae, *Grateloupia turuturu* Yamada, 1941 and *Asparagopsis armata* Harvey, 1855. Species unique to artificial habitats included the erect bryozoan, *T. inopinata*, the orange cloak sea squirt, *Botrylloides violaceus* Oka, 1927, an unidentified *Botrylloides* species, *Botrylloides* sp. indet. (Bishop et al., 2015b), the leathery sea squirt, *Styela clava* Herdman, 1881, the slipper limpet, *Crepidula fornicata* (Linnaeus, 1758) and the red alga, *Bonnemaisonia hamifera* Hariot, 1891. Semi-quantitative search-based techniques found 15 NNS across natural and artificial habitats, while quantitative quadrat techniques yielded only 8 species. The use of quantitative quadrat techniques alone failed to record *B. violaceus*, *C. fornicata*, *U. pinnatifida*, *G. turuturu*, *A. armata*, *B. hamifera* and the brown alga, *Sargassum muticum* (Yendo) Fensholt, 1955.

Statistical analysis of quantitative quadrat data found that mean NNS richness was significantly greater in artificial compared to natural habitats (Table 2a; Figure 2a). Similarly, assemblage composition varied significantly between natural and artificial habitats (Table 2a). Statistical analysis of semi-quantitative search-based data revealed that the mean richness did not differ significantly between natural and artificial habitats, although community assemblage did differ (Table 2b; Figure 2b, c). SIMPER analysis of quantitative quadrat data showed that over 80% of dissimilarity in assemblage composition between natural and artificial

habitats was attributed to *A. modestus* (32.6%), the red ripple bryozoan, *Watersipora subatra* (Ortmann, 1890) (30.9%) and the red alga, *Caulacanthus okamurae* Yamada, 1933 (23.1%), with all three species more abundant in artificial habitats. SIMPER analysis of semi-quantitative data revealed that > 50% of dissimilarity in assemblage composition between natural and artificial habitats was attributable to four species: *S. muticum* (17.3%), *W. subatra* (14.5%), the brown alga, *Colpomenia peregrina* Sauvageau, 1927 (12.4%) and the Pacific oyster, *Magallana gigas* (Thunberg, 1793) (11.8%). *S. muticum* and *C. peregrina* were more abundant in natural habitats, while *W. subatra* and *M. gigas* were more common in artificial habitats.

Table 2. PERMANOVA results comparing NNS richness and assemblage composition between natural and artificial habitats using (a) quantitative quadrat data and (b) semi-quantitative search-based data. Significant p-values are in bold.

(a) Quantitative data

Two-way PERMANOVA comparing species richness between natural and artificial habitats.

Source	df	SS	MS	Pseudo-F	P(perm)
Habitat	1	18976	18976	7.1199	0.0009
Site(Habitat)	18	47985	2665.8	33.327	0.0001
Residual	380	30396	79.989		
Total	399	97274			
Transformation:	pres/abs				

Two-way PERMANOVA comparing assemblage composition between natural and artificial habitats.

Source	df	SS	MS	Pseudo-F	P(perm)
Habitat	1	24297	24297	6.3912	0.0006
Site(Habitat)	18	68443	3802.4	16.724	0.0001
Residual	380	86397	227.36		
Total	399	179000			
Transformation:	fourth root				

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(b) Semi-quantitative data

One-way ANOVA comparing species richness between natural and artificial habitats.

Source	df	SS	MS	Pseudo-F	P(perm)
Habitat	1	94.912	94.912	0.59709	0.4781
Residual	19	3020.2	158.96		
Total	20	3115.1			
Transformation: pres/abs					

One-way PERMANOVA comparing assemblage composition between natural and artificial habitats.

Source	df	SS	MS	Pseudo-F	P(perm)
Habitat	1	4854.7	4854.7	9.041	0.0001
Residual	19	10202	536.96		
Total	20	15057			
Transformation: fourth root					

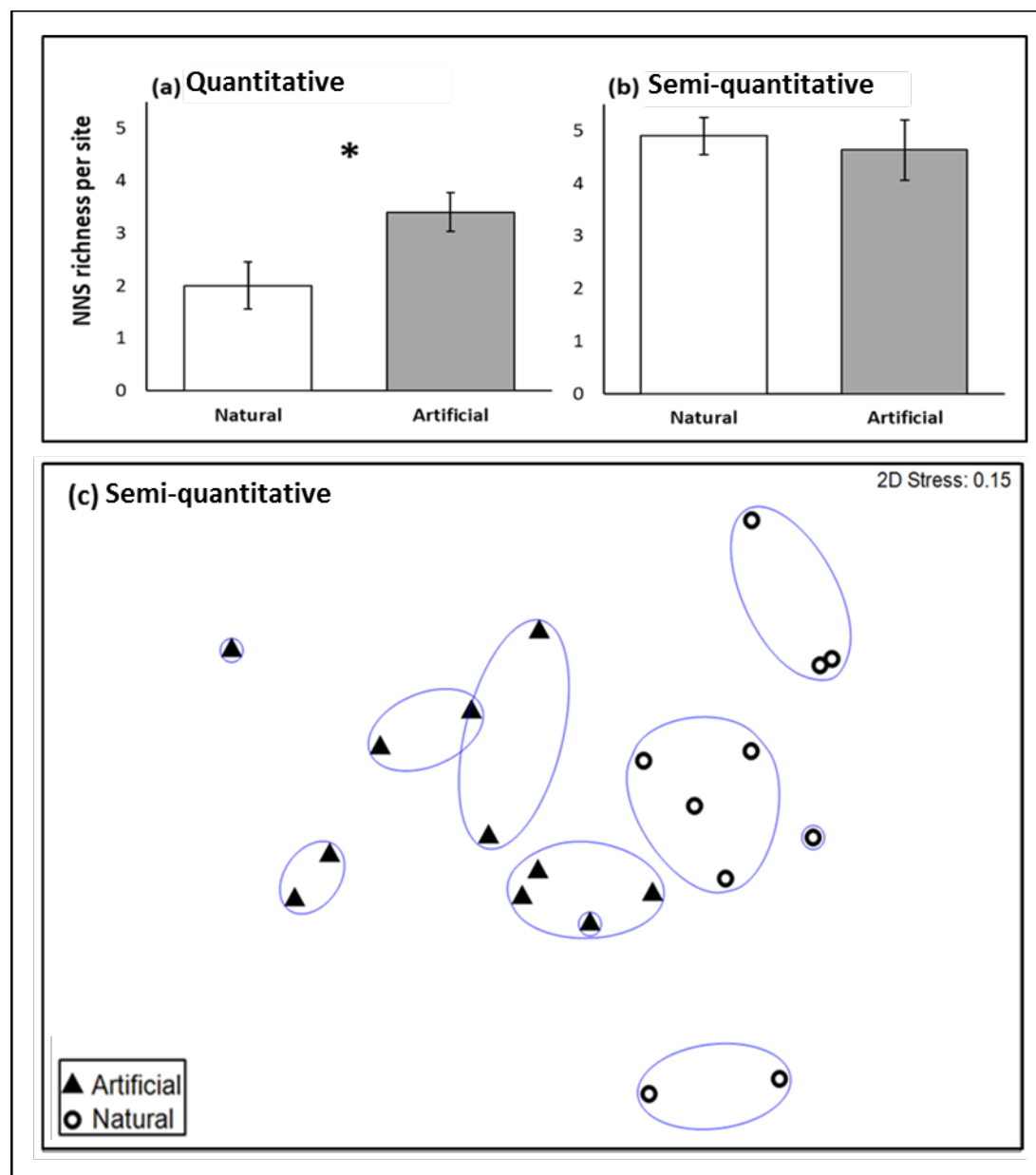


Figure 2. Comparison of mean number of NNS per site in natural and artificial intertidal habitats using (a) quantitative quadrat and (b) semi-quantitative search-based sampling techniques. *NNS richness was significantly greater in artificial compared to natural habitats using quantitative techniques ($p = 0.0009$) but not using semi-quantitative methods (natural sites, $n = 10$; artificial sites, $n = 11$). Error bars show standard error. (c) Non-metric multi-dimensional scaling plot (nMDS) showing significant difference in assemblage composition between natural and artificial sites using semi-quantitative search-based data. Assemblage composition between natural and artificial habitats differed significantly ($p = 0.0001$). The blue envelopes show a resemblance level of 75%.

3.3. Comparison of NNS in artificial habitats along the south coast of England

Overall, 26 NNS were recorded in artificial sites across eleven harbours. The barnacle, *A. modestus*, was encountered most (43 sites), while the erect bryozoan, *Bugulina simplex* (Hincks, 1886), the orange-tipped sea squirt, *Corella eumyota* Traustedt, 1882, *B. wrightii*, the green alga, *Codium fragile subsp. fragile* (Suringar) Hariot, 1889 and *U. pinnatifida* were only found at a single site each. Results are reported from both sampling methods, but only semi-quantitative search-based data were used to produce figures because this method captured more NNS overall.

Statistical analysis of quantitative quadrat data revealed that mean richness and assemblage composition were significantly different among harbours and regions (Table 3a), with the Central region supporting greater mean and total richness than the West ($p = 0.0222$) and East regions ($p = 0.0039$). Pairwise comparisons among harbours within regions revealed significantly higher richness in Torbay compared to Looe ($p = 0.0084$), Southampton compared to Portsmouth ($p = 0.0030$) and Shoreham compared to Folkestone ($p = 0.0045$). Pairwise comparisons showed that assemblage composition between the West and Central ($p = 0.0012$) and the Central and East ($p = 0.0012$) regions differed significantly. Differences in assemblage composition were found between Falmouth and Looe ($p = 0.0232$), Falmouth and Plymouth ($p = 0.0292$), Falmouth and Torbay ($p = 0.0244$), Looe and Torbay ($p = 0.0073$), Poole and Southampton ($p = 0.0090$), Southampton and Portsmouth ($p = 0.0002$) and Shoreham and Folkestone ($p = 0.0034$). SIMPER analysis revealed that the erect bryozoan, *Bugula neritina* (Linnaeus, 1758) contributed the most to the dissimilarity between West and Central (15%; greater in Central); whilst *W. subatra* contributed the most to dissimilarities between West and East (37.3%; greater in West) and Central and East regions (22.1%; greater in Central).

Analysis of semi-quantitative search-based data revealed a significant difference in NNS richness among regions, with the Central region supporting greater mean and total richness per harbour compared to West ($p = 0.0472$) and East regions ($p = 0.0014$; Table 3b, Figures

3, 4b). There were no significant differences, however, in mean richness among harbours within regions (Table 3b, Figure 4a). Assemblage composition varied significantly among both harbours and regions (Table 3b, Figure 5). Post-hoc pairwise tests comparing regions showed that the West and Central ($p = 0.0144$) and Central and East ($p = 0.0326$) assemblage compositions differed significantly. Comparisons of harbours within regions found that assemblage composition differed significantly between Falmouth and Plymouth ($p = 0.0355$), Poole and Portsmouth ($p = 0.0131$), Poole and Southampton ($p = 0.0014$) and Southampton and Portsmouth ($p = 0.0013$). SIMPER analysis revealed that *W. subatra* contributed the most to the dissimilarity between West and East regions (25.4%; greater in West), while *C. okamurae* contributed the most to dissimilarities between West and Central (11.4%; greater in West) and Central and East regions (1.6%; greater in East).

364 Table 3. PERMANOVA results for comparison of NNS richness and assemblage composition
365 using (a) quantitative quadrat data and (b) semi-quantitative search-based data in artificial
366 habitats among harbours and regions. Significant p-values are in bold.

(a) Quantitative data

Three-way PERMANOVA comparing species richness among regions, harbours and sites.

Source	df	SS	MS	Pseudo-F	P(perm)
Region	2	60707	30354	7.0537	0.0025
Harbour(Region)	7	118340	16905	3.9287	0.0019
Site(Harbour(Region))	30	118340	4303.2	15.623	0.0001
Residual	760	118340	275.44		
Total	799	118340			
Transformation:	pres/abs				

Three-way PERMANOVA comparing assemblage composition among regions, harbours and sites.

Source	df	SS	MS	Pseudo-F	P(perm)
Region	2	75905	37953	4.5881	0.0040
Harbour(Region)	7	221400	31628	3.8236	0.0001
Site(Harbour(Region))	30	248160	8272	11.922	0.0001
Residual	760	527330	693.85		
Total	799	1155800			
Transformation:	fourth root				

(b) Semi-quantitative data

Two-way PERMANOVA comparing species richness among regions and harbours.

Source	df	SS	MS	Pseudo-F	P(perm)
Region	2	4014	2007	7.0275	0.0012
Harbour(Region)	8	2399.2	299.9	1.0501	0.4183
Residual	33	9424.5	285.59		
Total	43	18170			
Transformation:	pres/abs				

Two-way PERMANOVA comparing assemblage composition among regions and harbours.

Source	df	SS	MS	Pseudo-F	P(perm)
Region	2	12842	6420.9	7.586	0.0001
Harbour(Region)	8	15168	1896	2.24	0.0028
Residual	33	27931	846.41		

This is an accepted proof.

O'Shaughnessy, K.A., Hawkins, S.J., Yunnice, A.L., Hanley, M.E., Lunt, P., Thompson, R.C. and Firth, L.B., 2020. Occurrence and assemblage composition of intertidal non-native species may be influenced by shipping patterns and artificial structures. *Marine Pollution Bulletin*, 154, 111082.
<https://doi.org/10.1016/j.marpolbul.2020.111082>

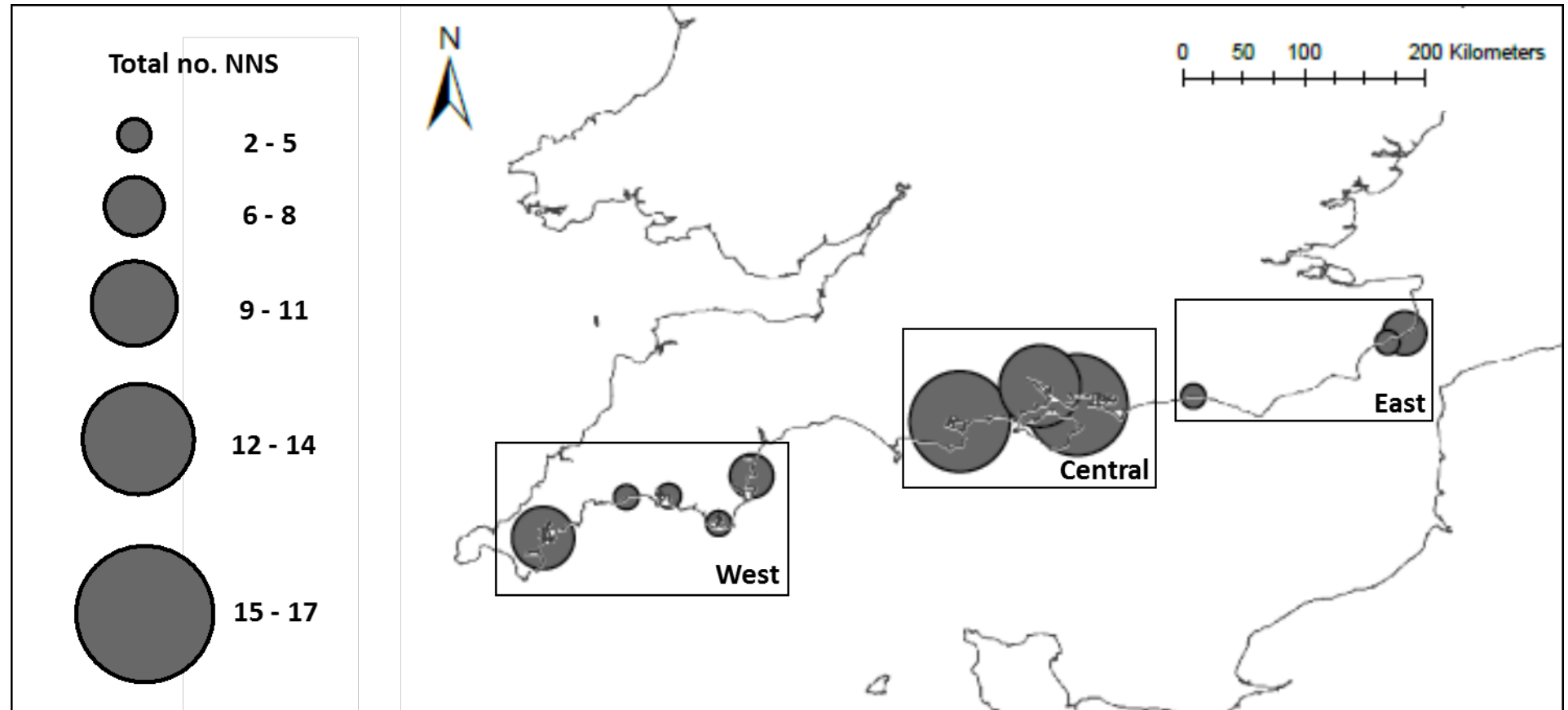
Total	43	65127
Transformation:	fourth root	

367

This is an accepted proof.

O'Shaughnessy, K.A., Hawkins, S.J., Yunnice, A.L., Hanley, M.E., Lunt, P., Thompson, R.C. and Firth, L.B., 2020. Occurrence and assemblage composition of intertidal non-native species may be influenced by shipping patterns and artificial structures. *Marine Pollution Bulletin*, 154, 111082.

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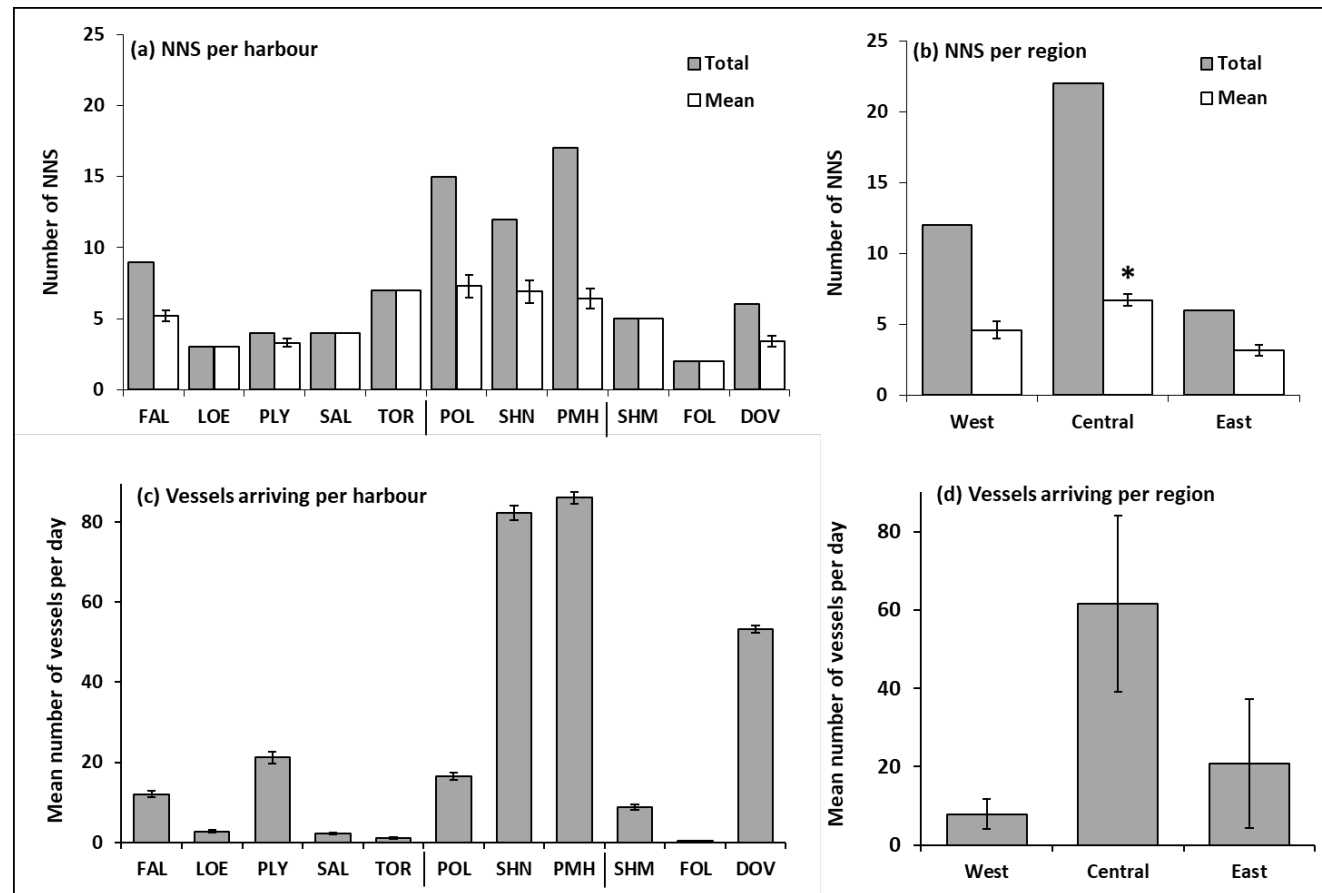
368

369 Figure 3. The number of NNS recorded per harbour from Rapid Assessment Surveys of intertidal artificial habitats along the south coast of
 370 England ranged from 2-17. Size of circles represents the total number of NNS recorded per harbour. Harbours from west to east: Falmouth,
 371 Looe, Plymouth, Salcombe, Torbay (West region), Poole, Southampton, Portsmouth (Central region), Shoreham, Folkestone and Dover (East
 372 region). Figure was produced using the semi-quantitative search-based data.

This is an accepted proof.

O'Shaughnessy, K.A., Hawkins, S.J., Yunnice, A.L., Hanley, M.E., Lunt, P., Thompson, R.C. and Firth, L.B., 2020. Occurrence and assemblage composition of intertidal non-native species may be influenced by shipping patterns and artificial structures. *Marine Pollution Bulletin*, 154, 111082.

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<https://doi.org/10.1016/j.marpolbul.2020.111082>

376 Figure 4. Total and mean number of NNS recorded from intertidal artificial habitats across the south coast of England per (a) harbour and (b)
377 region. *Numbers of NNS were significantly greater in Central compared to the West ($p = 0.0472$) and East ($p = 0.0014$) regions. Error bars
378 represent standard error. Bars showing means in (a) without a standard error bar represent harbours where only one site was surveyed. Mean
379 number of vessels arriving per day (c) by harbour and (d) by region were averaged over 60 days. Figure was produced using semi-quantitative
380 search-based data. Information was obtained from Marine Traffic (2019) using vessel data from February and March 2019.

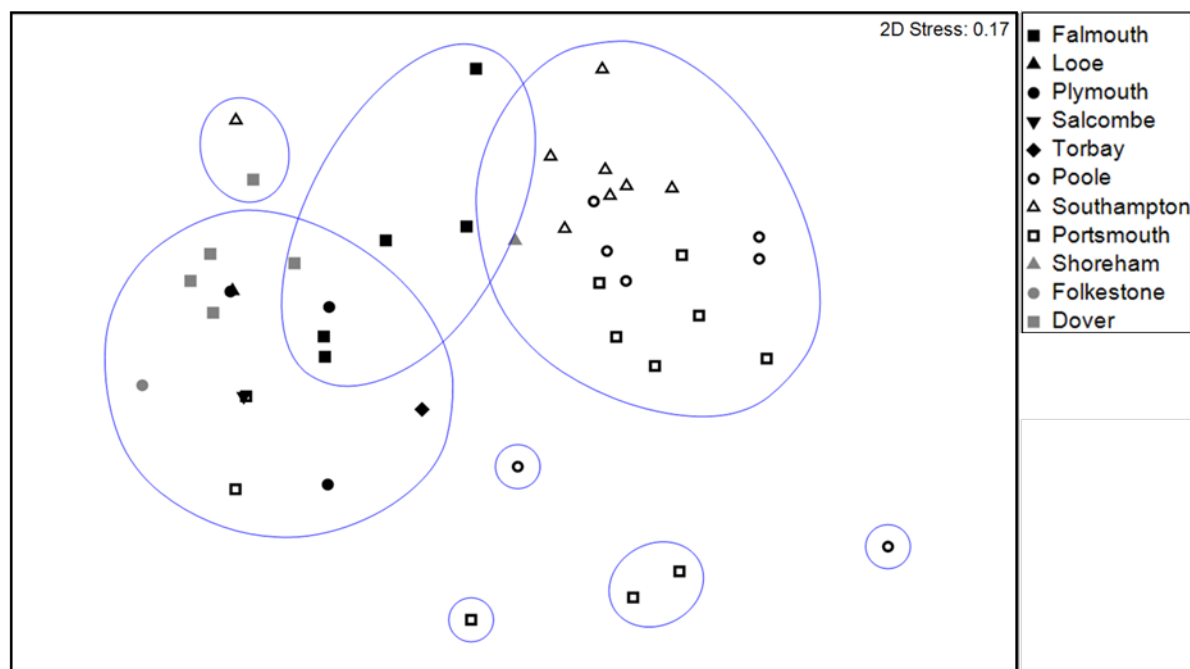


Figure 5. Non-metric multi-dimensional scaling plot (nMDS) showing significant variation in NNS assemblage composition from Rapid Assessment Surveys of intertidal artificial sites among harbours ($p = 0.0028$) and regions ($p = 0.0001$) along the south coast of England. Dark shapes represent harbours in the West region, open shapes indicate harbours in the Central region and grey shapes represent harbours in the East. Figure was created using semi-quantitative search-based data. The blue envelopes show a resemblance level of 60%.

3.4. Numbers and types of vessels by harbour

Portsmouth and Southampton supported the greatest average number of vessel arrivals per day at 86.4 and 83.1, respectively, while Folkestone supported the fewest (0.4; Figure 4c). The Central region supported the greatest average number of vessels per day at 61.6, while the West supported the fewest at 7.9 (Figure 4d). International shipping (container liner services) occurs out of Southampton with 12 carriers, as well as Plymouth and Dover with one carrier each (Table 1). Seven of the 12 container liner carriers that visit Southampton operate globally, with destinations in North and South America, Asia, India, the Pacific Islands, Australia, Europe, the Middle East, Africa and the Caribbean (World Port Source, 2019). The carrier that operates out of Plymouth is also a global carrier (similar destinations as listed

above), while the carrier out of Dover only operates out of northern Europe. International cruise liners operate out of Southampton with global destinations, while cruise liners out of Dover have European destinations only. Ferries to northern Europe (France and Spain) operate from Plymouth, Poole, Portsmouth and Dover, while military bases are located in Falmouth, Plymouth and Portsmouth (Table 1, Figure 1).

4. Discussion

Twenty-six non-native sessile invertebrates and macroalgae were found during Rapid Assessment Surveys of artificial structures in harbours along the south coast of England. Comparisons of NNS between natural and artificial habitats found that assemblage compositions differed significantly, but differences in richness depended on the sampling technique employed. The Central region supported greater mean and total richness, as well as different assemblage compositions, compared to the West and East regions. These differences might be attributed to regional shipping patterns, as most harbours in this study with high NNS richness saw relatively large amounts of vessel traffic.

Our study provided mixed evidence to support the hypothesis that artificial sites would support greater NNS richness compared to natural sites, as results differed depending on sampling method employed. Analysis of quantitative stratified-random quadrat data found differences in richness between natural and artificial habitats, while analysis of semi-quantitative search-based data (obtained from timed searches including complex habitats) did not detect differences. Both sampling techniques, however, showed that NNS assemblages between natural and artificial habitats were indeed different. These results agreed to some extent with previous studies that found assemblages of NNS differed between natural and artificial subtidal habitats (Glasby et al., 2007; Tyrrell and Byers, 2007; Dafforn et al., 2012). Conversely, the result that richness did not differ between natural and artificial habitats from

our study contrasts with work from these same studies listed above which showed that artificial supported more NNS than natural habitats. For example, Glasby et al. (2007) found that numbers of NNS were markedly greater on floating pontoons and pilings than on natural rocky reef. Furthermore, Tyrrell and Byers (2007) and Dafforn et al. (2012) experimentally showed that non-native fouling species outcompeted native species on artificial structures, but non-natives were not able to gain a foothold on natural substrate. Importantly, all of these studies were done in subtidal habitats (many on floating pontoons), which may explain the conflicting results. In the current study, semi-quantitative search-based sampling techniques allowed complex microhabitats to be searched (e.g., rock pools, crevices and gaps between boulders). Natural substrate generally provides more topographic complexity compared to artificial structures (Moschella et al., 2005; Chapman and Underwood, 2011). Thus, it is not surprising that many more NNS were observed during the searches in natural than artificial habitats. For example, whilst *S. muticum*, *C. pergrina*, *U. pinnatifida*, *A. armata* and *G. turuturu* were found in previous studies of artificial habitats (Arenas et al., 2006; Firth et al., 2013; Bishop et al., 2015b), they were only found in rock pools during timed searches in natural habitats in our study. This indicates the importance of water-retaining features for the successful establishment of non-native species on typically diverse natural rocky shores. These features have previously been identified as being important for providing shade and water retention at low tide to alleviate desiccation for native species in intertidal habitats (Firth et al., 2013, Firth et al., 2016). Conversely, in the subtidal zone, as desiccation is not a concern, topographic complexity may be needed for entirely different reasons, such as providing larvae and propagules refuge from predators or wave movement (Kovalenko et al., 2012; Strain et al., 2017). Different uses of topographic complexity by resident organisms between intertidal and subtidal habitats may be a reason for the differential results obtained between previous surveys and the current survey.

The frequency of arriving vessels, types and departure points of vessels arriving in harbours, as well as the subsequent secondary destinations to which species can be transported (Carlton, 1996b; Ruiz et al., 2000; Clarke Murray et al., 2011), may explain the high number and variety of NNS around Southampton and the Central region in general. The success of an introduced species is highly dependent on propagule pressure (Lockwood et al., 2005; Copp et al., 2010), and thus it is not surprising that most of the harbours supporting relatively high species richness were also harbours that supported a high frequency of arriving vessels. For example, Southampton was unique in our study in being one of the largest cruise ship and container ports in the UK, and the busiest along the English south coast (Associated British Ports, 2019; Marine Traffic, 2019; World Port Source, 2019). As such, it sees significantly more international traffic than any other of the south coast ports. Global movement of NNS into the British Isles has been described by Eno et al. (1997) who proposed scenarios depicting likely invasion pathways. Many species were proposed to have been transferred directly to the British Isles from their places of origin, such as *B. hamifera* from Southeast Asia, *A. modestus* from Australia and *C. fornicata* from the eastern seaboard of the US. These routes of transfer explain high propagule pressure in the port of Southampton, where cargo tankers regularly arrive from international ports. However, Eno et al. (1997) suggested many other non-natives (i.e., *A. armata*, *S. muticum*, *C. peregrina*) were first transported from their origin to continental Europe, followed by a secondary transfer into the British Isles across the English Channel. Bishop et al. (2015a) provided evidence for this, as their Rapid Assessment Surveys showed a general pattern of northward movement of NNS from Brittany, France across the English Channel to South West England over time. This invasion pathway might explain the high numbers of NNS in Portsmouth (and to a lesser degree, Poole), where large passenger ferries arrive from northern European destinations. The current study can only suggest the above as invasion pathways, as identifying NNS from source ports was outside the scope of this study. Thus, future research can focus on providing evidence by ground-

truthing these results or using molecular markers to show that the same NNS occurred in the departing international harbours as the receiving ports in southern England.

Although the relationship between NNS richness and number of vessels arriving per day was clear at the regional level (i.e., West, Central, East), at the harbour level, there were some harbours that did not follow this trend. Poole Harbour supported a relatively high number of NNS, yet the number of arriving vessels was relatively low compared to other harbours with high species richness (i.e., Portsmouth and Southampton). This suggests that factors other than propagule pressure from international shipping play a role in determining success of NNS. The geomorphology of Poole is quite different than the other harbours in this study, in that the harbour itself has a very narrow mouth, a double high tide daily and the smallest tidal range in the study area (1.8 m), as well as very poor flushing (Humphreys, 2005; May, 2005; World Port Source, 2019). Poole has been described as a lagoon-like harbour; it is a shallow and warm body of water (Humphreys, 2005; May, 2005). These conditions may be particularly favourable for NNS originating from warmer waters, such as *D. vexillum* (Zaiko et al., 2007; Lambert, 2009). Moreover, slow and incomplete flushing of the harbour means larvae of NNS are present in the water for long periods of time, potentially allowing increased settlement compared to harbours with faster flushing times. These conditions are also favourable to yachting, and so smaller leisure craft regularly travel among harbours within the Solent (Central region; SJH pers. obs.), potentially acting as secondary mobile vectors (Clarke Murray et al., 2011). On the other hand, Dover sees a relatively high number of vessels per day but supports relatively few numbers of NNS. Although the numbers of vessels are high in Dover, the origins and destinations are almost exclusively European (Marine Traffic, 2019; World Port Source, 2019). This effectively means lower propagule pressure from global invaders directly. Moreover, the geomorphology of Dover is different from the other harbours in this study, in that Dover is not a natural bay or inlet. Rather, the port was artificially created when the Dover Southern Breakwater was constructed at the beginning of the 20th century. It

is therefore likely that the hydrodynamics – which can affect turbidity and scouring of structures (Govarets and Lauwaert, 2009; Dugan et al., 2011), and can dictate transport of larvae in currents and tides (McQuaid and Phillips, 2000) – differ markedly between Dover and naturally-formed ports.

Evidence from this study and others suggest that artificial structures probably play an important role in the initial establishment and then secondary transport of NNS away from their initial point of introduction (Eno et al., 1997; Neves et al., 2007; Mineur et al., 2012). By nature, docks and floating pontoons are constantly in close proximity to cargo tankers, passenger ferries and recreational vessels, and thus there is a high probability of species spreading from mobile vectors (vessels) to stationary structures (floating pontoons, docks; Neves et al., 2007). In a study examining the fouling communities of boat hulls and associated floating pontoons and concrete structures in an international Brazilian port, Neves et al. (2007) found that biotic communities on hulls were similar to those on the pontoons; while communities on concrete structures were a similar but smaller subset of the species found on hulls. This is because boat hulls and floating pontoons rise and fall with the tide, while concrete structures are fixed in place (similar to intertidal natural rocky shores). Hulls of recreational boats are regularly cleaned of fouling organisms (Neves et al., 2007); it is even becoming increasingly common for transoceanic vessels to undergo regular hull cleaning (Hopkins and Forrest, 2008; PML Applications Ltd, 2019). Thus, the biological communities they support are typically younger (i.e. at earlier successional stages) than those on floating pontoons and associated docks. As such, pontoons act as “reservoirs” of established NNS communities (Neves et al., 2007; Floerl et al., 2009; Foster et al., 2016), while concrete structures support fewer numbers of species, but of which have the ability to invade intertidal natural habitats (Neves et al., 2007; Epstein and Smale, 2018). In our study, harbours east of Torbay were dominated by sedimentary substrata, thus fouling organisms typical of natural rocky shores arriving from distant hard substrata have only been able to establish and survive by colonising artificial structures in

these otherwise uninhabitable areas. In this way, artificial structures may affect ecological connectivity by providing “stepping stones” for the movement of species across seascapes (Sammarco et al., 2004; Airoidi et al., 2015; Bishop et al., 2017). Moreover, smaller harbours (e.g., Folkestone) with many fewer artificial structures compared to larger harbours (e.g., Portsmouth) provided less hard substrata for NNS spread from a mobile vector to a stationary structure. These smaller harbours also supported less diverse NNS communities due to lower diversity of artificial structures (e.g., Folkestone Harbour is largely composed of rock and concrete seawalls and lacks floating pontoons). As such, these smaller harbours are less likely to act as NNS reservoirs.

The limitations of sampling natural and artificial sites equally are extremely challenging to overcome; yet a combined sampling approach (quantitative stratified-random quadrats and semi-quantitative timed searches), like those employed in this study, can address some of the problems associated with sampling in these habitats. By nature, complex microhabitats on gently sloping natural rocky shores (e.g., rock pools, gaps between boulders) are generally easier (and safer) to sample compared to those on artificial structures. This was acknowledged in the methods of the current study by slightly extending the timed search on structures that were difficult to sample due to safety and logistical reasons. For example, boulder groynes and riprap revetment provide internal compartments created by the stacking of boulders to maximise coastal protection. The interior of these structures provide functional niches that are absent on the exterior that protect organisms from desiccation, wave exposure and sand scour; thus species diversity tends to be greater within the internal compartments, which are difficult to access/observe (Sherrard et al., 2016; Liversage and Chapman, 2018). Traditional quadrat sampling avoids gaps between boulders and other complex microhabitats, consequently missing vital species diversity information. Therefore, numbers and abundances of NNS recorded from the exterior of these structures probably do not accurately represent the true NNS diversity of the entire structure. On the other hand, limited areal extent provided

by other artificial structures, such as seawalls, means that the likelihood of these structures to be fully sampled ("censused"; Chapman et al., 2018) using quadrat sampling techniques is much greater than on a large natural rocky shore or more complex artificial structures, where many diverse habitats are likely to be missed (Chapman et al., 2018). In the current study, quadrat sampling was able to cover most of the available area on seawalls but missed areas in larger natural sites (e.g., rock pools). If quantitative quadrat sampling alone had been used (which is a typical method in biodiversity surveys), our study would have concluded that intertidal artificial habitats support greater (mean) numbers of NNS (per unit area) than natural habitats. By employing a timed search, this study came to a very different conclusion - that of which challenges the commonly accepted concept that artificial structures support greater NNS richness compared to natural rocky shores. Most NNS surveys along the south coast of England have focused on surveying marina pontoons in subtidal habitats (Arenas et al., 2006; Ashton et al., 2006; Bishop et al., 2015a,b; Foster et al., 2016), as these are known "hot spots" for introduction. Our study demonstrated that areas not labelled as "hot spots", such as natural rocky shores, should not be ignored. A recent report cautioned that natural rocky shores might facilitate "spillover" of NNS from populations in marinas to natural habitats, facilitating the spread out from the initial sites of introduction (Epstein and Smale, 2018). Although not explicitly tested for, our study showed that intertidal natural sites do indeed support many NNS and may actually contribute to their spread between major transport hubs and surrounding bays and harbours.

Results from this study could be strengthened by formal tests on the effects of local environmental and physical conditions on success of NNS establishment. Physical factors, such as hydrodynamics (Horvath and Crane, 2010; Zardi et al., 2006) and pollution load (Dafforn et al., 2011; McKenzie et al., 2011) are important in influencing NNS colonisation and competition with native biota. For example, previous studies have shown that industrial and urban runoff adversely affects native composition and ecological functioning of marine

communities (Johnston and Roberts, 2009; Burton and Johnston, 2010), and that species response (recolonisation) can vary depending on exposure to particular contaminants (Trannum et al., 2004). Additionally, NNS have been shown to tolerate contaminants and pollution while abundances of native species decline under the same conditions (Crooks et al., 2011; Dafforn et al., 2011; McKenzie et al., 2011). Thus, a decline in native biotic communities can easily give NNS a competitive advantage where conditions are unfavourable for native species (Johnston et al., 2017).

The loss of natural habitat caused by coastal development and ocean sprawl is leading to the need to explore alternative options to traditional hard built structures for coastal protection (Dafforn et al., 2015a; Firth et al., 2016a). There is therefore an increasing impetus to ecologically enhance hard structures to fulfil secondary management goals, such as increase biodiversity, enhance ecosystem services or reduce abundance of NNS (i.e., "ecological engineering"; Dafforn, 2017; Evans et al., 2017; Strain et al., 2019a,b). Results from our surveys clearly demonstrate that ecological engineering designs must consider the potential for colonisation by NNS (Sella and Perkol-Finkel, 2015; Dafforn, 2017; Strain et al., 2017). Traditionally, ecological engineering interventions that have included rock pools retrofitted onto seawalls (Chapman and Underwood, 2011; Browne and Chapman, 2014) or drilling pits into seawalls or breakwaters (Firth et al., 2014; Evans et al., 2016), have been advocated as a means of enhancing species diversity through water retention. Our survey showed that these interventions may increase the risk of colonisation by NNS, such as *S. muticum* and *U. pinnatifida*, which were regularly found in rock pools in natural sites. Information from biodiversity studies and Rapid Assessment Surveys should thus serve as a benchmark against which to measure change to biotic communities over time, and is an essential first step in informing management decisions concerning design details for ecological engineering of artificial structures in coastal intertidal habitats (Dafforn et al., 2015b; Mayer-Pinto et al., 2017).

Ocean sprawl is accelerating the rate of NNS introduction and spread, contributing to biotic homogenisation and the growing biodiversity crisis. Understanding the ecological role of artificial structures in the marine and coastal environments is critical for preserving native biodiversity and building resilience to establishment of NNS. Our results suggest that global shipping and artificial structures may play an important role in the introduction and spread of NNS. Other factors such as local environmental conditions and geomorphology of harbours undoubtedly contribute to NNS success, but disentangling these factors is difficult. Therefore, all potential mechanisms of NNS introduction, establishment and spread need investigation so that ecologists might develop the predictive capability to identify areas at high risk of invasion, which can aid in effective forecasting for potential invaders.

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